

DESCRIPTION

COMPOSITION AND METHOD FOR PROMOTING HAIR GROWTH

TECHNICAL FIELD

The present invention relates to a composition
5 and method for promoting hair growth in a mammalian subject.

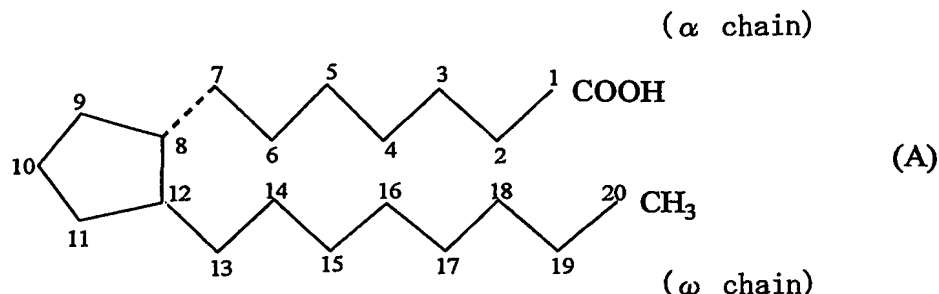
BACKGROUND ART

Hair loss or alopecia may result from genetic
factors, aging, local or systemic disease or certain
therapeutic drugs designed to alleviate conditions such as
10 cancer. Various preparations for preventing or reducing
hair loss and/or promoting hair growth are proposed, for
example those containing female hormones which can promote
blood circulation, reinforce hair root function, moisturize
scalp and inhibit male hormone function; 5 α -reductase
15 inhibitors; or minoxidil, trichosaccharides or the like as
main ingredients. However, they cannot show satisfactory
hair growth-promoting effects, and some may raise side-
effect problems such as sexual function disorder.

It is strongly desired to develop a hair growth-
20 promoting agent having superior effects without side effect.

Prostaglandins (hereinafter, referred to as
PG(s)) are members of class of organic carboxylic acids,
which are contained in tissues or organs of human or other
mammals, and exhibit a wide range of physiological activity.
25 PGs found in nature (primary PGs) generally have a

prostanoic acid skeleton as shown in the formula (A):



On the other hand, some of synthetic analogues of primary PGs have modified skeletons. The primary PGs are classified to PGAs, PGBs, PGCs, PGDs, PGEs, PGFs, PGGs, PGHs, PGIs and PGJs according to the structure of the five-membered ring moiety, and further classified into the following three types by the number and position of the unsaturated bond at the carbon chain moiety:

Subscript 1: 13,14-unsaturated-15-OH

Subscript 2: 5,6- and 13,14-diunsaturated-15-OH

Subscript 3: 5,6-, 13,14-, and 17,18-triunsaturated-15-OH.

Further, the PGFs are classified, according to the configuration of the hydroxyl group at the 9- and 11-position, into α type (the hydroxyl group is of an α-configuration) and β type (the hydroxyl group is of a β-configuration).

Certain prostaglandin compound having two hetero atoms at the 15 position are known in the art. US patent No. 4,088,775 discloses certain 15-ethylenedioxy-prostanoic

acid. In addition, US patent No. 4,870,104 discloses 11
halo prostaglandins which may have an
ethylenedioxymethylene group at 15-position and use thereof
as agents inhibiting gastric acid secretion. Further, US
5 patent No. 6,353,014 discloses a certain 15-ketal analogs
of F series prostaglandins useful for treating ocular
hypertension and glaucoma.

Those prior arts do not disclose nor suggest that
prostaglandin compound having two hetero atoms at the 15
10 position may be useful in the stimulation of hair growth.

DISCLOSURE OF THE INVENTION

An object of the present invention is to provide
a composition for promoting hair growth in a mammalian
subject.

15 Further object of the present invention is to
provide a method for promoting hair growth in a mammalian
subject.

Still further object of the present invention is
to provide a novel compound useful for promoting hair
20 growth in a mammalian subject.

Namely, the present invention relates to a
composition for promoting hair growth in a mammalian
subject which comprises a prostaglandin compound having two
hetero atoms at the 15 position as an active ingredient
25 thereof.

Further, the present invention relates to a method for promoting hair growth in a mammalian subject, which comprises topically administering a prostaglandin compound having two hetero atoms at the 15 position to the
5 subject in need thereof.

Furthermore, the present invention relates to use of a prostaglandin compound having two hetero atoms at the 15 position for manufacturing a composition for promoting hair growth in a mammalian subject.

10 Still further, the present invention relates to a novel prostaglandin compound having two hetero atoms at the 15 position.

DETAILED DESCRIPTION OF THE INVENTION

The nomenclature of the PG compounds used herein
15 is based on the numbering system of the prostanoic acid represented in the above formula (A).

The formula (A) shows a basic skeleton of the C-20 carbon atoms, but the present invention is not limited to those having the same number of carbon atoms. In the
20 formula (A), the numbering of the carbon atoms which constitute the basic skeleton of the PG compounds starts at the carboxylic acid (numbered 1), and carbon atoms in the α -chain are numbered 2 to 7 towards the five-membered ring, those in the ring are 8 to 12, and those in the ω -chain are
25 13 to 20. When the number of carbon atoms is decreased in

the α -chain, the number is deleted in the order starting from position 2; and when the number of carbon atoms is increased in the α -chain, compounds are named as substitution compounds having respective substituents at position 2 in place of the carboxy group (C-1). Similarly, when the number of carbon atoms is decreased in the ω -chain, the number is deleted in the order starting from position 20; and when the number of carbon atoms is increased in the ω -chain, the carbon atoms beyond position 20 are named as substituents. Stereochemistry of the compounds is the same as that of the above formula (A) unless otherwise specified.

In general, each of the terms PGD, PGE and PGF represents a PG compound having hydroxy groups at positions 9 and/or 11, but in the present specification, these terms also include those having substituents other than the hydroxy group at positions 9 and/or 11. Such compounds are referred to as 9-dehydroxy- 9-substituted-PG compounds or 11-dehydroxy-11-substituted-PG compounds. A PG compound having hydrogen in place of the hydroxy group is simply named as 9- or 11-dehydroxy-PG compound.

As stated above, the nomenclature of the PG compounds is based on the prostanoic acid skeleton. However, in case the compound has a similar partial structure as a prostaglandin, the abbreviation of "PG" may be used. Thus, a PG compound of which α -chain is extended

by two carbon atoms, that is, having 9 carbon atoms in the α -chain is named as 2-decarboxy-2-(2-carboxyethyl)-PG compound. Similarly, a PG compound having 11 carbon atoms in the α -chain is named as 2-decarboxy-2-(4-carboxybutyl)-PG compound. Further, a PG compound of which ω -chain is extended by two carbon atoms, that is, having 10 carbon atoms in the ω -chain is named as 20-ethyl-PG compound. These compounds, however, may also be named according to the IUPAC nomenclatures.

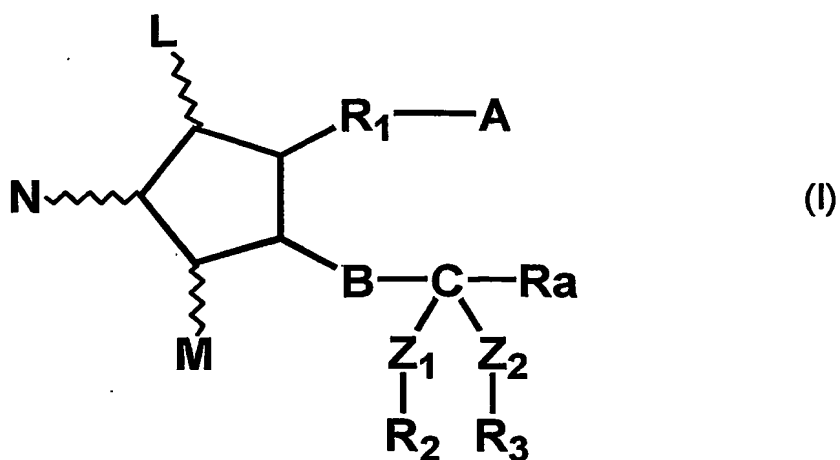
Examples of the analogs (including substituted derivatives) or derivatives include a PG compound of which carboxyl group at the end of α -chain is esterified; a compound of which α -chain is extended; physiologically acceptable salt thereof; a compound having a double bond at 2-3 position or a triple bond at position 5-6, a compound having substituent(s) at position 3, 5, 6, 16, 17, 18, 19 and/or 20; and a compound having lower alkyl or a hydroxy (lower) alkyl group at position 9 and/or 11 in place of the hydroxy group.

According to the present invention, preferred substituents at position 3, 17, 18 and/or 19 include alkyl having 1-4 carbon atoms, especially methyl and ethyl. Preferred substituents at position 16 include lower alkyl such as methyl and ethyl, hydroxy, halogen atoms such as chlorine and fluorine, and aryloxy such as

trifluoromethylphenoxy. Preferred substituents at position 17 include lower alkyl such as methyl and ethyl, hydroxy, halogen atoms such as chlorine and fluorine, aryloxy such as trifluoromethylphenoxy. Preferred substituents at position 20 include saturated or unsaturated lower alkyl such as C1-4 alkyl, lower alkoxy such as C1-4 alkoxy, and lower alkoxy alkyl such as C1-4 alkoxy-C1-4 alkyl. Preferred substituents at position 5 include halogen atoms such as chlorine and fluorine. Preferred substituents at position 6 include an oxo group forming a carbonyl group. Stereochemistry of PGs having hydroxy, lower alkyl or hydroxy(lower)alkyl substituent at position 9 and/or 11 may be α , β or a mixture thereof.

Further, the above analogs or derivatives may be compounds having an alkoxy, cycloalkyl, cycloalkyloxy, phenoxy or phenyl group at the end of the ω -chain where the chain is shorter than the primary PGs.

A preferred prostaglandin compound used in the present invention is represented by formula (I):



wherein L, M and N are hydrogen, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo, wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have at least one double bond;

A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

B is $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$;

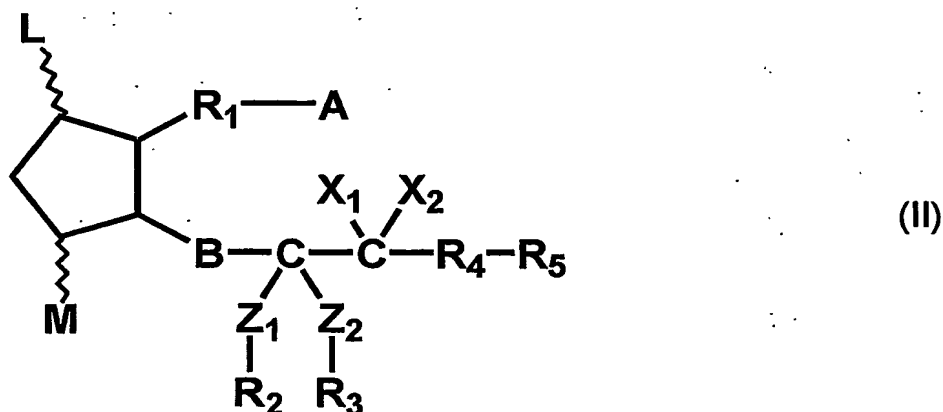
Z₁ and Z₂ are oxygen, nitrogen or sulfur,

R₂ and R₃ are optionally substituted lower alkyl, which is optionally linked together to form lower alkylene,

R₁ is a saturated or unsaturated bivalent lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur; and

Ra is a saturated or unsaturated lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, oxo, hydroxy, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group; lower alkoxy; lower alkanoyloxy; cyclo(lower)alkyl; cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; heterocyclic-oxy group.

A more preferred prostaglandin compound used in the present invention is represented by the formula (II):



wherein L and M are hydrogen, hydroxy, halogen, lower alkyl, hydroxy(lower)alkyl, lower alkanoyloxy or oxo, wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have one or more double bonds;

A is $-\text{CH}_3$, or $-\text{CH}_2\text{OH}$, $-\text{COCH}_2\text{OH}$, $-\text{COOH}$ or a functional derivative thereof;

B is $-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-$ or $-\text{C}\equiv\text{C}-$;

Z_1 and Z_2 are oxygen, nitrogen or sulfur,

R_2 and R_3 are optionally substituted lower alkyl, which is optionally linked together to form lower alkylene,

X_1 and X_2 are hydrogen, lower alkyl, or halogen;

5 R_1 is a saturated or unsaturated bivalent lower or medium aliphatic hydrocarbon residue, which is unsubstituted or substituted with halogen, alkyl, hydroxy, oxo, aryl or heterocyclic group, and at least one of carbon atom in the aliphatic hydrocarbon is optionally substituted
10 by oxygen, nitrogen or sulfur;

R_4 is a single bond or lower alkylene; and

R_5 is lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group.

15 In the above formula, the term "unsaturated" in the definitions for R_1 and R_4 is intended to include at least one or more double bonds and/or triple bonds that are isolatedly, separately or serially present between carbon atoms of the main and/or side chains. According to the
20 usual nomenclature, an unsaturated bond between two serial positions is represented by denoting the lower number of the two positions, and an unsaturated bond between two distal positions is represented by denoting both of the positions.

25 The term "lower or medium aliphatic hydrocarbon"

refers to a straight or branched chain hydrocarbon group having 1 to 14 carbon atoms (for a side chain, 1 to 3 carbon atoms are preferable) and preferably 1 to 10, especially 1 to 8 carbon atoms.

5 The term "halogen atom" covers fluorine, chlorine, bromine and iodine.

The term "lower" throughout the specification is intended to include a group having 1 to 6 carbon atoms unless otherwise specified.

10 The term "lower alkyl" refers to a straight or branched chain saturated hydrocarbon group containing 1 to 6 carbon atoms and includes, for example, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, t-butyl, pentyl and hexyl.

15 The term "lower alkylene" refers to a straight or branched chain bivalent saturated hydrocarbon group containing 1 to 6 carbon atoms and includes, for example, methylene, ethylene, propylene, isopropylene, butylene, isobutylene, t-butylene, pentylene and hexylene.

20 The term "lower alkoxy" refers to a group of lower alkyl-O-, wherein lower alkyl is as defined above.

The term "hydroxy(lower)alkyl" refers to a lower alkyl as defined above which is substituted with at least one hydroxy group such as hydroxymethyl, 1-hydroxyethyl, 2-hydroxyethyl and 1-methyl-1-hydroxyethyl.

25

The term "lower alkanoyloxy" refers to a group represented by the formula RCO-O- , wherein RCO- is an acyl group formed by oxidation of a lower alkyl group as defined above, such as acetyl.

5 The term "cyclo(lower)alkyl" refers to a cyclic group formed by cyclization of a lower alkyl group as defined above but contains three or more carbon atoms, and includes, for example, cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

10 The term "cyclo(lower)alkyloxy" refers to the group of cyclo(lower)alkyl-O-, wherein cyclo(lower)alkyl is as defined above.

 The term "aryl" may include unsubstituted or substituted aromatic hydrocarbon rings (preferably
15 monocyclic groups), for example, phenyl, tolyl, xylyl. Examples of the substituents are halogen atom and halo(lower)alkyl, wherein halogen atom and lower alkyl are as defined above.

 The term "aryloxy" refers to a group represented
20 by the formula ArO- , wherein Ar is aryl as defined above.

 The term "heterocyclic group" may include mono- to tri-cyclic, preferably monocyclic heterocyclic group which is 5 to 14, preferably 5 to 10 membered ring having optionally substituted carbon atom and 1 to 4, preferably 1
25 to 3 of 1 or 2 type of hetero atoms selected from nitrogen

atom, oxygen atom and sulfur atom. Examples of the heterocyclic group include furyl, thienyl, pyrrolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, imidazolyl, pyrazolyl, furazanyl, pyranyl, pyridyl, pyridazinyl, 5 pyrimidyl, pyrazinyl, 2-pyrrolinyl, pyrrolidinyl, 2-imidazolinyl, imidazolidinyl, 2-pyrazolinyl, pyrazolidinyl, piperidino, piperazinyl, morpholino, indolyl, benzothienyl, quinolyl, isoquinolyl, purinyl, quinazolinyl, carbazolyl, acridinyl, phenanthridinyl, benzimidazolyl, 10 benzimidazolinyl, benzothiazolyl, phenothiazinyl. Examples of the substituent in this case include halogen, and halogen substituted lower alkyl group, wherein halogen atom and lower alkyl group are as described above.

The term "heterocyclic-oxy group" means a group 15 represented by the formula HcO- , wherein Hc is a heterocyclic group as described above.

The term "functional derivative" of A includes salts (preferably pharmaceutically acceptable salts), ethers, esters and amides.

20 Suitable "pharmaceutically acceptable salts" include conventionally used non-toxic salts, for example a salt with an inorganic base such as an alkali metal salt (such as sodium salt and potassium salt), an alkaline earth metal salt (such as calcium salt and magnesium salt), an 25 ammonium salt; or a salt with an organic base, for example,

an amine salt (such as methylamine salt, dimethylamine salt, cyclohexylamine salt, benzylamine salt, piperidine salt, ethylenediamine salt, ethanolamine salt, diethanolamine salt, triethanolamine salt, tris(hydroxymethylamino)ethane salt, monomethyl- monoethanolamine salt, procaine salt and 5 caffeine salt), a basic amino acid salt (such as arginine salt and lysine salt), tetraalkyl ammonium salt and the like.. These salts may be prepared by a conventional process, for example from the corresponding acid and base 10 or by salt interchange.

Examples of the ethers include alkyl ethers, for example, lower alkyl ethers such as methyl ether, ethyl ether, propyl ether, isopropyl ether, butyl ether, isobutyl ether, t-butyl ether, pentyl ether and 1-cyclopropyl ethyl 15 ether; and medium or higher alkyl ethers such as octyl ether, diethylhexyl ether, lauryl ether and cetyl ether; unsaturated ethers such as oleyl ether and linolenyl ether; lower alkenyl ethers such as vinyl ether, allyl ether; lower alkynyl ethers such as ethynyl ether and propynyl 20 ether; hydroxy(lower)alkyl ethers such as hydroxyethyl ether and hydroxyisopropyl ether; lower alkoxy (lower)alkyl ethers such as methoxymethyl ether and 1-methoxyethyl ether; optionally substituted aryl ethers such as phenyl ether, tosyl ether, t-butylphenyl ether, salicyl ether, 25 3,4-di-methoxyphenyl ether and benzamidophenyl ether; and

aryl(lower)alkyl ethers such as benzyl ether, trityl ether and benzhydryl ether.

Examples of the esters include aliphatic esters, for example, lower alkyl esters such as methyl ester, ethyl ester, propyl ester, isopropyl ester, butyl ester, isobutyl ester, t-butyl ester, pentyl ester and 1-cyclopropylethyl ester; lower alkenyl esters such as vinyl ester and allyl ester; lower alkynyl esters such as ethynyl ester and propynyl ester; hydroxy(lower)alkyl ester such as hydroxyethyl ester; lower alkoxy (lower) alkyl esters such as methoxymethyl ester and 1-methoxyethyl ester; and optionally substituted aryl esters such as, for example, phenyl ester, tolyl ester, t-butylphenyl ester, salicyl ester, 3,4-di-methoxyphenyl ester and benzamidophenyl ester; and aryl(lower)alkyl ester such as benzyl ester, trityl ester and benzhydryl ester.

The amide of A mean a group represented by the formula $\text{-CONR}'\text{R}''$, wherein each of R' and R'' is hydrogen, lower alkyl, aryl, alkyl- or aryl-sulfonyl, lower alkenyl and lower alkynyl, and include for example lower alkyl amides such as methylamide, ethylamide, dimethylamide and diethylamide; arylamides such as anilide and toluidide; and alkyl- or aryl-sulfonylamides such as methylsulfonylamide, ethylsulfonyl-amide and tolylsulfonylamide.

Preferred examples of L and M include hydroxy and

oxo, and especially, M and L are hydroxy groups which has a 5-membered ring structure of, so called, PGF type.

Preferred example of A is -COOH, its pharmaceutically acceptable salt, ester or amide thereof.

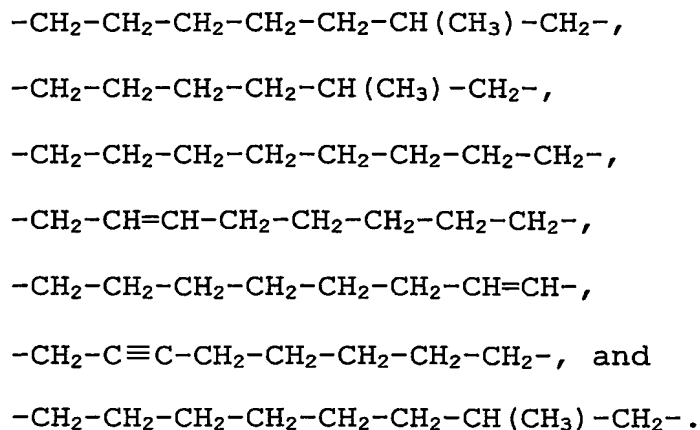
5 Preferred B is -CH₂-CH₂-, so called 13,14-dihydro type.

Preferred example of X₁ and X₂ is fluorine, so called 16,16-difluoro type.

10 Preferred R₁ is a hydrocarbon residue containing 1-10 carbon atoms, preferably 6-10 carbon atoms. Further, at least one carbon atom in the aliphatic hydrocarbon is optionally substituted by oxygen, nitrogen or sulfur.

Examples of R₁ include, for example, the following groups:

15 -CH₂-CH₂-CH₂-CH₂-CH₂-CH₂-,
-CH₂-CH=CH-CH₂-CH₂-CH₂-,
-CH₂-CH₂-CH₂-CH₂-CH=CH-,
-CH₂-C≡C-CH₂-CH₂-CH₂-,
-CH₂-CH₂-CH₂-CH₂-O-CH₂-,
20 -CH₂-CH=CH-CH₂-O-CH₂-,
-CH₂-C≡C-CH₂-O-CH₂-,
-CH₂-CH₂-CH₂-CH₂-CH₂-CH₂-CH₂-,
-CH₂-CH=CH-CH₂-CH₂-CH₂-CH₂-,
-CH₂-CH₂-CH₂-CH₂-CH₂-CH=CH-,
25 -CH₂-C≡C-CH₂-CH₂-CH₂-CH₂-,



Preferred Ra is a hydrocarbon containing 1-10 carbon atoms, more preferably, 1-8 carbon atoms. Ra may have one or two side chains having one carbon atom.

Preferred Z₁ and Z₂ are oxygen.

R₂ and R₃ are preferably linked together to form C2 or C3 alkylene.

The configuration of the ring and the α- and/or ω chains in the above formula (I) and (II) may be the same as or different from that of the primary PGs. However, the present invention also includes a mixture of a compound having a primary type configuration and a compound of a non-primary type configuration.

In the present invention, any of isomers such as the individual tautomeric isomers, the mixture thereof, or optical isomers, the mixture thereof, a racemic mixture, and other steric isomers may be used in the same purpose.

According to the present invention, a composition for promoting hair growth comprising the prostaglandin

compound defined as above as an active ingredient is applied to a mammalian subject in need of promotion of hair growth.

5 The term "hair" in the present specification and claims covers any hair on a mammalian subject, especially a human subject, for example, hairs on the top of the head, on the armpits, on the pubic area, on the face including eyelash, eyebrow, eyelid, mustache, beard and whisker, on the chest, arms and legs.

10 The term "promoting hair growth" in the present specification and claims covers not only promoting hair growth but also promoting hair germination and thickening hairs. As is shown in the examples below, the composition of the present invention has the effect of thickening the growing hair in addition to the promotion of hair growth.

15 According to the present invention, the composition may be provided as, for example, pharmaceuticals, quasi-drugs (i.e. iyakubugaihin in Japanese) or cosmetics. The composition may topically be applied for the purpose of promoting hair growth onto the surface of skin where hair growth is desired, such as scalp, face, beard, head, pubic area, upper lip, eyelash, eyebrow, and eyelid.

20 The dose of the prostaglandin compound in the composition of the present invention may vary according to

the compound to be used, the type of subject, age, skin area to be applied the composition, progress of baldness or desired effect, administration volume and period for treatment. Although a suitable concentration may be chosen
5 as desired, in a typical case wherein the composition is topically administrated to an adult, the formulation containing 0.0000001% - 10%, preferably 0.00001% - 5%, more preferably 0.0001% - 1% and especially 0.001-0.1% of the active ingredient can be applied 1-6 times, preferably 1-4
10 times per day.

The dosage form of the composition of the present invention can be any of known topically applicable forms. For example, but not limited thereto, lotion, tonic, emulsion, external drug creams such as liniments and milky
15 lotions, external semi-solid preparations such as ointments, paste, jelly and sprays. The composition may also be formulated as hair shampoos or hair rinses.

The composition of the present invention may further contain physiologically acceptable additives. Said
20 additives may include the ingredients used with the present compounds such as excipient, diluent, filler, resolvent, lubricant, adjuvant, binder, disintegrator, emulsifier, dispersing agent, suspending agent, thickener, tonicity agent, buffering agent, soothing agent, preservative,
25 antioxidant, corrigent, flavor, colorant, a functional

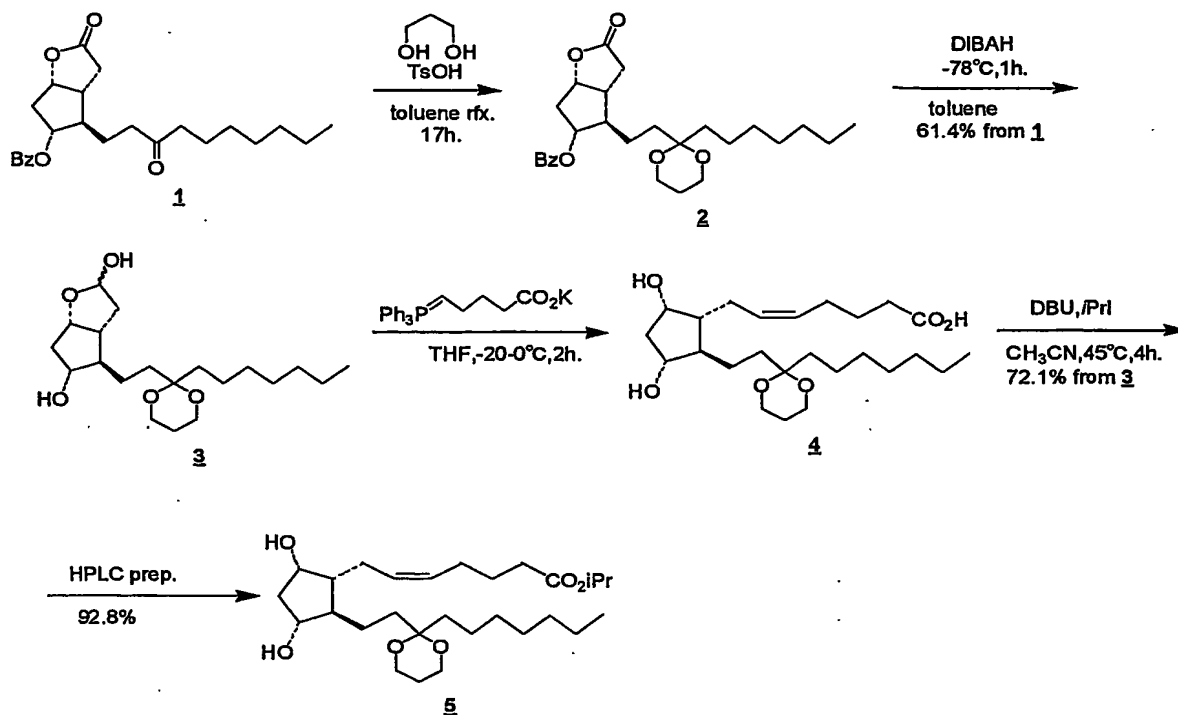
material such as cyclodextrin and biodegradable polymer, stabilizer. They may be further dissolved in an appropriate solvent such as fatty acid or its mono, di or triglyceride. The additives are well known to the art and
5 may be selected from those described in general reference books of pharmaceuticals or cosmetics.

The composition of the present invention may further contain other ingredients as far as they do not contradict the purpose of the present invention. The
10 composition may be prepared in a conventional manner for manufacturing the desired formulations by adding the prostaglandin compound defined as above.

The further details of the present invention will follow with reference to test examples, which, however, are
15 not intended to limit the present invention.

Synthesis example 1

13,14-dihydro-15,15-trimethylenedioxy-20-ethyl-PGF_{2α}
isopropyl ester (5)



To the solution of compound 1 (510.0mg, 1.273mmol) in toluene (10.2ml), 1,3-propanediol (0.92ml, 12.73mmol) and a catalytic amount of p-toluene sulfonic acid were added and the mixture was heated for 17 hours under reflux. After that, the reaction was left stood until it was cooled to room temperature, and washed with saturated aqueous sodium bicarbonate and saturated aqueous sodium chloride. The organic phase was dried with magnesium sulfate and evaporated under reduced pressure. The residue was purified by means of silica gel column chromatography (Merck 7734, Hexane: ethyl acetate=3:2) to give compound 2 (581.3mg).

The solution of compound 2 (580.0mg, 1.265mmol) in toluene (11.6ml) was cooled to -78°C, 1.5M-DIBALH (in

toluene, 2.95ml, 4.427mmol) was added dropwise thereto and the mixture was stirred for 1 hour, and then, methanol (1.79ml) was added dropwise to the resulting mixture. Saturated aqueous Rochelle salt (100ml) was added thereto and the mixture was vigorously stirred for 30 minutes. The resulting mixture was extracted with ethyl acetate, and the organic layer was washed with saturated salt water, dried with magnesium sulfate and evaporated under reduced pressure. The residue was purified by means of silica gel column chromatography (Merck 7734, Hexane: ethyl acetate=1:9-0:10) to give compound 3 (275.2mg, yield 61.4% from 1).

To the dispersion of (4-carboxybutyl)triphenyl phosphonium bromide (1.346g, 3.038mmol) in THF (6ml), 1M-potassium t-butoxide in THF (6.07ml, 6.07mmol) at 0°C was added. The reaction was stirred for 1 hour at room temperature and then cooled to -20°C. Compound 3 (269.2mg, 0.7594mmol) in THF (7ml) was added dropwise thereto and stirred for 2 hours at -20-0°C. Ice cold water was added to the reaction, THF was removed by evaporation evaporated under reduced pressure. To the concentrated residue at 0°C, ice cold 1N aqueous hydrochloric acid was added dropwise to adjust the solution to pH 4.

The solution was extracted with ethyl acetate and the organic layer was washed with saturated aqueous sodium

chloride, dried with magnesium sulfate and evaporated under reduced pressure. The residue was added with ether and stirred for 17 hours at room temperature and then, filtrated with celite. The filtrate was evaporated under reduced pressure to give crude compound 4.

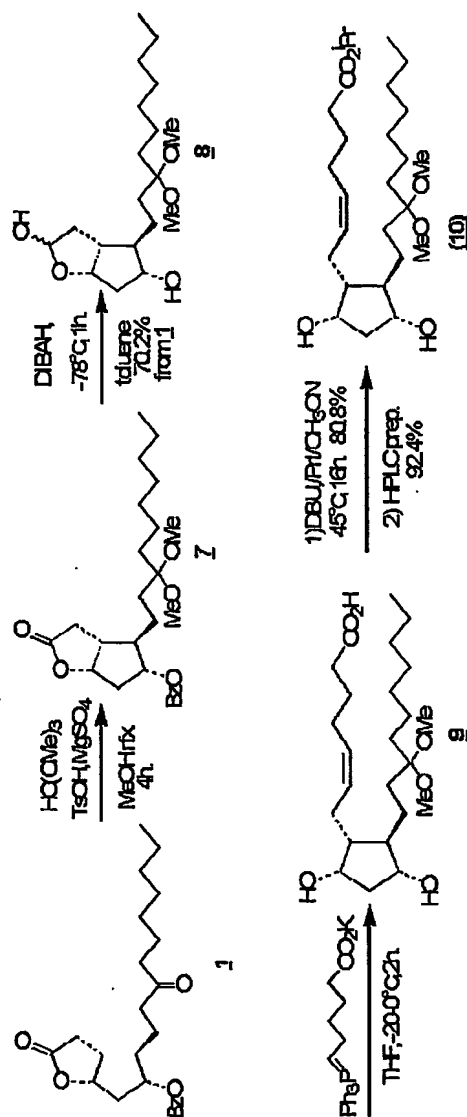
Compound 4 (0.7594mmol) in acetonitrile (7.6ml) was added with DBU (0.45ml, 3.038mmol), isopropyl iodide (0.30ml, 3.038 mmol) and stirred for 4 hours at 45°C. The reaction mixture was evaporated under reduced pressure. The residue was added with water and extracted with ethyl acetate. The organic layer was washed with saturated aqueous sodium chloride solution, dried with magnesium sulfate and evaporated under reduced pressure. The residue was purified by means of silica gel column chromatography (Merck 9385, hexane : ethyl acetate= 2:3) to give 727.2mg of the desired product (yield 72.1% from 3). Thus obtained compound 4 (carboxylic acid, 259.0mg) was further purified by separation HPLC to give compound 5 (isopropyl ester, 240.3mg, HPLC purification yield 92.8%).

¹H-NMR spectrum (200MHz, CDCl₃) of compound 5: δ 5.57-5.14 (2H, m), 5.01 (1H, sept, J=6.2Hz), 4.17 (1H, bs), 3.97 (1H, bs), 4.00-3.78 (4H, m), 2.76 (1H, d, J=6.2Hz), 2.29 (2H, t, J=7.5Hz), 2.44-2.06 (5H, m), 1.88 (2H, bt), 1.93-1.18 (22H, m), 1.23 (6H, d, J=6.2Hz), 0.89 (3H, t, J=6.8Hz)

Synthesis example 2

13,14-dihydro-15,15-dimethoxy-20-ethyl-PGF_{2α} ester (10)

isopropyl



5

To the solution of compound 1 (797.8mg, 2.002mmol) in methanol (2.4ml), a catalytic amount of p-toluene sulfate, methyl orthoformate (2.19ml, 20.02mmol) and unhydrous magnesium sulfate (1.20g, 10.01mmol) were added

and heated under reflux for 4 hours. The reaction was cooled and added with sodium hydrogen carbonate, and filtered with celite. The filtrate was evaporated under reduced pressure and the residue was purified by means of silica gel column chromatography (Merck 7734g, hexane : ethyl acetate = 3:2) to give compound 7 (884.3mg, yield 98.9%).

The solution of compound 7 (767.5mg, 1.719mmol) in toluene(15.4ml) was cooled to -78°C, 1.5M-DIBAH (in toluene, 4.0ml, 6.016mmol) was added dropwise thereto and the mixture was stirred for 1 hour. Then, methanol was added dropwise to the reaction and the reaction was heated to room temperature. Saturated aqueous Rochelle salt (150ml) was added thereto and the mixture was vigorously stirred for 30 minutes. The resulting mixture was extracted with ethyl acetate, and the organic layer was washed with saturated salt water, dried with magnesium sulfate and evaporated under reduced pressure. The residue was purified by means of silica gel column chromatography (Merck 9385, hexane: ethyl acetate=1:9) to give compound 8 (415.8mg, yield 70.2%).

To the dispersion of (4-carboxybuthyl)triphenyl phosphonium bromide (1.250g, 2.819mmol) in THF, 1M-potassium t-butoxide in THF (5.64ml, 5.64mmol) at 0 °C was added. The reaction was stirred for 1 hour at room

temperature and then cooled to -20°C . Compound 8 (242.8mg, 0.7048mmol) in THF (4ml) was added dropwise thereto and stirred for 2 hours at $-20-0^{\circ}\text{C}$. Ice cold water was added to the reaction, and THF was removed by evaporation under reduced pressure. To the residue at 0°C , ice cold 1N aqueous hydrochloric acid was added dropwise to adjust the solution to pH 5. The solution was extracted with ethyl acetate and the organic layer was washed with saturated aqueous sodium chloride, dried with magnesium sulfate and evaporated under reduced pressure. The residue was added with ether and stirred for 17 hours at room temperature and then, filtrated with celite. The filtrate was evaporated under reduced pressure to give crude compound 9 (carboxylic acid).

To the solution of compound 9 (0.7048mmol) in acetonitrile (7ml), DBU (0.42ml, 2.819mmol), isopropyl iodide (0.28ml, 2.819mmol) were added and the mixture was stirred for 16 hours at 45°C . The reaction mixture was evaporated under reduced pressure. The residue was added with water and extracted with ethyl acetate. The organic layer was washed with saturated aqueous sodium chloride, dried with magnesium sulfate and evaporated under reduced pressure. The residue was purified by means of silica gel column (Merck 9385, hexane:ethyl acetate= 1:2) to give compound 10 (268.0mg, yield 80.8% from 8).

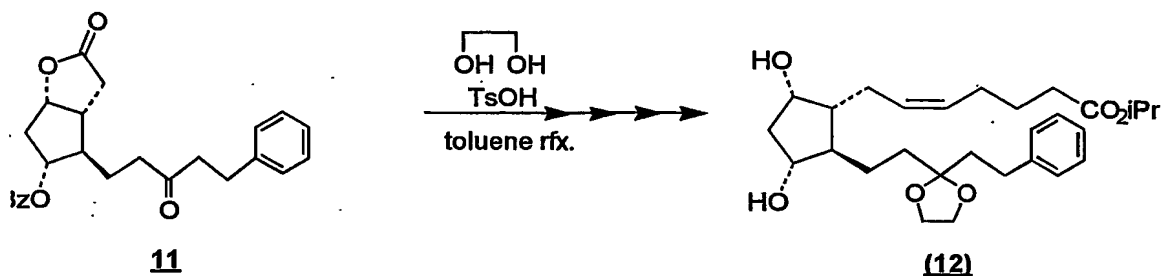
Compound 10 obtained as above (total 370 mg) was further purified by separation HPLC to give purified compound 10 (341.9mg, HPLC purification yield 92.4%).

¹H-NMR spectrum (200MHz, CDCl₃) of compound 10: δ5.54-5.13(2H, m), 5.00(1H, sept, J=6.2Hz), 4.18(1H, bs), 3.95(1H, bs), 3.16(6H, s), 2.66(1H, d, J=6.4Hz), 2.29(2H, t, J=7.3Hz), 2.48-2.06(5H, m), 1.89(2H, bt), 1.79-1.17(20H, m), 1.23(6H, d, J=6.2Hz), 0.89(3H, t, J=6.8Hz)

Synthesis example 3

13,14-dihydro-15,15-ethylenedioxy-17-phenyl-18,19,20-trinor-PGF_{2α} isopropyl ester (12)

Compound 12 was prepared from compound 11 in a same manner as Synthesis example 1.



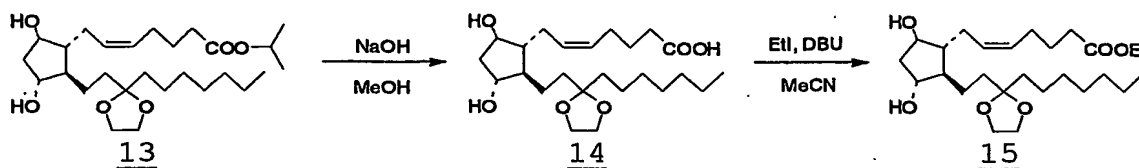
¹H-NMR spectrum (200MHz, CDCl₃) of compound 11: δ8.04-7.93(2H, m), 7.63-7.38(3H, m), 7.35-7.11(5H, m), 5.21-5.03(2H, m), 2.98-2.24(11H, m), 2.12-1.98(1H, m), 1.80-1.50(2H, m)

¹H-NMR spectrum (200MHz, CDCl₃) of compound 12: δ7.35-7.12(5H, m), 5.56-5.35(2H, m), 5.00(1H, sept, J=6.2Hz), 4.15(1H, bs), 3.96(4H, s), 3.92(1H, bs), 3.18(1H, bd),

2.86(1H, bd), 2.75-2.63(2H, m), 2.28(2H, t, J=7.3Hz), 2.46-1.15(17H, m), 1.22(6H, d, J=6.2Hz)

Synthesis example 4

13,14-dihydro-15,15-ethylenedioxy-20-ethyl-PGF2 α ethyl ester (15)



To the solution of compound 13 (9.18 g, 19.59 mmol) in methanol (91.8 ml), 8N-aqueous sodium hydroxide (24.49 ml) was added at 0°C. The reaction mixture was stirred for 3 hours at room temperature, and then acidified with 6N-hydrochloric acid at 0°C. The mixture was extracted with ethyl acetate (100 ml + 50 ml). The organic layer was washed with saturated aqueous sodium chloride (100 ml x 2), dried over anhydrous magnesium sulfate. The extract was evaporated under reduced pressure to obtain crude acid 14 as oil.

To the solution of crude acid 14 and 1,8-diazabicyclo[5.4.0]undec-7-ene (11.72 ml) in acetonitrile (60 ml), ethyl iodide (6.27 ml) was added dropwise at 0°C. The reaction mixture was stirred at 45°C for 17 hours, then cooled to room temperature, and evaporated. To the residue, water (100 ml) was added. The mixture was extracted with ethyl acetate (100 ml x 2). The organic layer was washed

with 0.1N-hydrochloric acid, saturated aqueous sodium bicarbonate (100 ml) and saturated aqueous sodium chloride (100 ml). The extract was dried over anhydrous magnesium sulfate and evaporated. The residue was purified by two
5 times of silica gel column chromatography (Merck 7734, 220 g, hexane : ethyl acetate= 2:3, -> BW-300, 210g, hexane:2-Propanol=6:1) to obtain ethyl ester 15 (8.60 g, 18.92 mmol, ,96.6% from 13) as a colorless oil.

¹H-NMR (200MHz in CDCl₃, TMS = 0ppm) of the compound 15: δ
10 5.58-5.29(2H, m), 4.15(1H, brs), 4.13(2H, q, J=7.1Hz), 3.97(1H, brs), 3.94(4H, s), 2.80-2.70(1H, br), 2.49-2.36(1H, m), 2.32(2H, t, J=7.4Hz), 2.36-2.15(4H, m), 1.90-1.83(2H, m), 1.83-1.12(20H, m), 1.26(3H, t, J=7.1Hz), 0.88(3H, t, J=6.5Hz)

15 Example 1

Eight-weeks-old male C3H/HeN mice were used. The hair on the back was clipped by electric clipper so that the hair in the clipped area was removed as much as possible. Three days after the clipping, mice without
20 visible scratches were selected and used in this study. Each group consisted of 3 animals. After group assignment, the groups were housed separately in aluminum cages (3 animals/cage, 180 mm W x 300 mm D x 130 mm H; Nippon Cage, Ltd., Japan).

25 Each test compound was dissolved in 70% (w/w)

aqueous ethanol. Each dose formulation of test compound was evenly applied topically once daily (100 μ L per mouse) to the clipped dorsal skin area (approximately 2 \times 4 cm), except for Saturday and Sunday, for 30 days. The control group received an equal amount of the vehicle in the same manner.

Macroscopic observations of the hair growth were performed 14, 16, 18, 21, 23, 25, 28 and 30 days after the start of the treatment. Hair growth was scored according to the scale below:

- no hair growth observed
- \pm hair growth \leq 10% of the clipped area
- + hair growth 10 - 40% of the clipped area
- ++ hair growth 40 - 80% of the clipped area
- +++ hair growth \geq 80% of the clipped area

The results are shown in Table 1. In the vehicle-treated control group, no hair growth was observed during the treatment period. In the 0.01% and 0.1% compound A-treated groups, a dose-dependent hair growth was noted, and all animals treated with 0.1% compound A showed a hair growth classified to the highest score (+++) at Days 28 and 30 of treatment. In the 0.1% compound B-treated group, hair growth score for 2 out of 3 animals were +++ and that of the remaining animal was ++ at the end of the treatment period. In the 0.1% compound C-treated group, hair growth

was observed on 1 out of 3 animals. In the 0.1% compound D-treated group, hair growth was observed on 2 out of 3 animals

Table 1. Effects of Topical Application of Compound A, B, C and D on Hair Growth in C3H/HeN Mice

Groups	AnimalNo.	Hair Growth Score							
		Days of Treatments							
		14	16	18	21	23	25	28	30
Control (Vehicle)	0101	-	-	-	-	-	-	-	-
	0102	-	-	-	-	-	-	-	-
	0103	-	-	-	-	-	-	-	-
Compound A 0.1%	0301	-	±	±	+	++	+++	+++	+++
	0302	-	-	±	±	++	++	+++	+++
	0303	-	±	±	±	++	++	+++	+++
Compound A 0.01%	0201	-	-	-	-	-	-	±	±
	0202	-	-	-	-	-	+	+	+
	0203	-	-	±	±	±	+	++	++
Compound B 0.1%	0501	-	-	±	±	+	++	++	++
	0502	-	-	-	±	+	+	++	+++
	0503	-	-	-	±	+	++	++	+++
Compound C 0.1%	0701	-	-	-	-	-	-	-	-
	0702	-	-	±	±	±	+	+	+
	0703	-	-	-	-	-	-	-	-
Compound D 0.1%	0901	-	-	-	-	-	-	-	-
	0902	-	±	±	+	+	+	++	++
	0903	-	-	-	-	-	-	±	±

Compound A: 13,14-dihydro-15,15-ethylenedioxy-20-ethyl-PGF_{2α} isopropyl ester

Compound B: 13,14-dihydro-15,15-ethylenedioxy-17-phenyl-18,19,20-trinor-PGF_{2α} isopropyl ester

Compound C: 13,14-dihydro-15,15-trimethylenedioxy-20-ethyl-

PGF_{2α} isopropyl ester

Compound D: 13,14-dihydro-15,15-dimethoxy-20-ethyl-PGF_{2α}
isopropyl ester

Example 2

5 Eight-weeks-old male C3H/HeN mice were used. The
hair on the back was clipped by electric clipper so that
the hair in the clipped area was removed as much as
possible. Three days after the clipping, mice without
visible scratches were selected and used in this study.
10 Each group consisted of 3 animals. After group assignment,
the groups were housed separately in aluminum cages (3
animals/cage, 180 mm W x 300 mm D x 130 mm H; Nippon Cage,
Ltd., Japan).

15 Test compound was dissolved in 75% aqueous
ethanol. The formulation of test compound (compound E) was
evenly applied topically once daily (100 μL per mouse) to
the clipped dorsal skin area (approximately 2 x 4 cm),
except for Saturday and Sunday, for 23 days. The control
group received an equal amount of the vehicle in the same
20 manner.

Macroscopic observations of the hair growth were
performed 14, 16, 18, 21 and 23 days after the start of the
treatment. Hair growth was scored according to the scale
shown as above. Results are shown in Table 2. In the 0.1%
25 compound E-treated group, hair growth was observed on 3 out

of 3 animals.

Table 2. Effects of Topical Application of Compound E on Hair Growth in C3H/HeN Mice

Groups	AnimalNo.	Hair Growth Score				
		Days of Treatments				
		14	16	18	21	23
Control (Vehicle)	0101	-	-	-	-	-
	0102	-	-	-	-	-
	0103	-	-	-	-	-
Compound E 0.1%	0201	-	-	±	±	+
	0202	-	-	±	++	++
	0203	-	-	-	+	++

Compound E: 13,14-dihydro-15,15-ethylenedioxy-20-ethyl-

5 PGF_{2α} ethyl ester

Example 3

31 days after the start of the treatment conducted in Example 1, grown-up hairs in the treated area and hairs in the untreated area (i.e. non clipped area) were collected, respectively. Enlarged photomicrographs of the collected hairs were taken. Thickness of the randomly selected each ten hairs were measured and calculated the average.

Results are shown in Table 3. The results show that the hair grown-up by treating with the specific prostaglandin of the present invention was thicker than

those in the untreated area.

Table 3. Effects of Topical Application of Compounds A and B on Thickness of Grown-up Hair in C3H/HeN Mice

Hair group	n	Hair thickness, μm mean \pm SE
Control area	3	27.8 \pm 2.6
0.1% Compound A-treated area	3	34.1 \pm 1.6
Control area	3	29.3 \pm 1.0
0.1% Compound B-treated area	3	33.3 \pm 0.7